10/505291

Sertifikaat ENTKANTOOR REPUBLIC OF SOUTH AFRICA

DEPARTEMENT VAN HANDEL EN NYWERHEID



PCT/IB 03 /_0 0 5 0 0

Certificate

PATENT OFFICE REPUBLIEK VAN SUID-AFRIKA

> DEPARTMENT OF TRADE AND INDUSTRY

> > 19 AUG 2004

Hiermee word gesertifiseer dat This is to certify that

REC'D 0 8 APR 2003

WIPO PCT

the documents attached hereto are true copies of the Forms P2, P6, provisional specification and drawings of South African Patent Application No. 2002/1395 in the name of ADCOCK INGRAM LIMITED

Filed

19 February 2002

Entitled

PHARMACEUTICAL

COMPOSITIONS

PRIORIT

SUBMITTED OR TRANSMITTED IN COMPLIANCE WITH RULE 17.1(a) OR (b)

Geteken te

in die Republiek van Suid-Afrika, hierdie PRETORIA in the Republic of South Africa, this

28th

dag van day of

February 2003

Registrateur van Patente Registrar of Patents

PUBLIC OF SOUTH AFRICA REGISTER OF PATENTS					PATENTS ACT, 1978		
OFFICIAL APPLICATION LODGING DATE: PROVISIONAL ACCEPTANCE D.					EPTANCE DATE		
21 01 2002/1393	22	22 19 FEB 2002			2	47	
INTERNATIONAL CLASSIFICATION	LOD	LODGING DATE: COMPLETE				GRA	NTED DATE
51	23						
FULL NAME(S) OF APPLICANT(S)/PATENTEE	(S)			•			
71 ADCOCK INGRAM LIMITED							
APPLICANTS SUBSTITUTED:					•	DAT	E REGISTERED
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ASSIGNEE(S)						D/	ATE REGISTERED
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FULL NAME(S) OF INVENTOR(S)							
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72 TO BE ADVISED PRIORITY CLAIMED COUNTRY	• •		NUM	BER		DAT	E
	NIL		NUM	BER NI	L	DAT	≣ NIL
PRIORITY CLAIMED COUNTRY N.B. Use International abbreviation for country (see Schedule 4) 33	NIL				L		
PRIORITY CLAIMED COUNTRY N.B. Use International abbreviation for country 33			31		L		
PRIORITY CLAIMED COUNTRY N.B. Use International abbreviation for country (see Schedule 4) TITLE OF INVENTION 54 PHARMACEUTICAL COMPOSI	TIONS		31		L		
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REPUBLIC OF SOUTH AFRICA REPUBLIC OF SOUTH AFRICA REPUBLIC OF SOUTH AFRICA PATENTS ACT, 1978

APPLICATION FOR A PATENT
AND ACKNOWLEDGEMENT OF RECEIP 19.02.02
(Section 30 (1) - Regulation 22)

R0060,00

The gr	ranting of a patent is hereby requested by the undermentioned appl	licant on the basis of the present application filed in duplicate
	OFFICIAL APPLICATION NO.	REPUBLIEK YAN SULD AFRIKA
21	01,2002/1395	PA132630/P
	FULL NAME(S) OF APP	PLICANT(S)
71	ADCOCK INGRAM LIMITED	
	ADDRESS(ES) OF APE	PLICANT(S)

17 HARRISON AVENUE, BRYANSTON, GAUTENG, SOUTH AFRICA

TITLE OF INVENTION

PHARMACEUTICAL COMPOSITIONS

THE APPLICANT CLAIMS PRIORITY AS SET OUT ON THE ACCOMPANYING FORM P.2. THE EARLIEST PRIORITY CLAIM IS:

COUNTRY: NIL NUMBER: NIL DATE: NIL

THIS APPLICATION IS FOR A PATENT OF ADDITION TO PATENT APPLICATION NO.

THIS APPLICATION IS A FRESH APPLICATION IN TERMS OF SECTION 37 AND IS BASED ON APPLICATION NO.

21 01

THIS APPLICATION IS ACCOMPANIED BY:

\boxtimes	 A single copy of a provisional specification of 5 pages.
	2. Drawings of sheets
	Publication particulars and abstract (Form P.8 in duplicate).
	A copy of Figure of the drawings (if any) for the abstract.
	5. Assignment of Invention.
	6. Certified priority document.
	7. Translation of the priority document.
	8. Assignment of priority rights.
	9. A copy of the Form P.2 and the specification of S.A. Patent Application No
	10. Declaration and power of attorney on Form P.3.
	11. Request for ante-dating on Form P.4.
	12. Request for classification on Form P.9.
×	13. Form P.2 in duplicate.

74 ADDRESS FOR SERVICE: SPOOR & FISHER, SANDTON

Dated: 19 February 2002

14. Other.

SPOOR & FISHER
PATENT ATTORNEYS FOR THE APPLICANT(S)

RECEIVED

REGISTRAR OF PATENTS DESIGNS,

TRADE MARKS AND COPYRIGHT

2002 -02- 19

REGIS: RATEUR VAN PATENTE, MODELLE, HANDETSMERKBÆM OMFFLINSREG



PROVISIONAL SPECIFICATION

(Section 30(1) - Regulation 27)

	OFFICIAL APPLICATION NO.		LODGING DATE			
21	01 2002/1395	22	19 FEBRUARY 2002			
	FULL NAMES OF AP	PLICANTS				
71	ADCOCK INGRAM LIMITED		•			
L						
	5.11. MAY 50.05 M	n/CNTOD				
	FULL NAMES OF IN	VENTOR				
72	TO BE ADVISED		•			
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TITLE OF INVENTION						
	11125, 31					
54	PHARMACEUTICAL COMPOSITIONS	•				

BACKGROUND OF THE INVENTION

This invention relates to pharmaceutical compositions and their use in the symptomatic relief and treatment of pain, with or without fever.

SUMMARY OF THE INVENTION

According to one aspect of the invention, a pharmaceutical composition comprises a combination of (i) an analgesic, (ii) a selective or specific COX-2 inhibitor, and (iii) an opiate, and a pharmaceutically acceptable carrier.

In a preferred composition of the invention the analgesic (i) is paracetamol or a pharmaceutically acceptable salt or derivative thereof, the selective or specific COX-2 inhibitor (ii) is selected from the group comprising meloxicam, celecoxib, rofecoxib and pharmaceutically acceptable salts or derivatives thereof, and the opiate (iii) is selected from the group

comprising codeine, morphine, tramadol, fentanyl and pharmaceutically acceptable salts or derivatives thereof.

A particularly preferred composition of the invention comprises a combination of paracetamol, meloxicam and codeine phosphate.

The invention extends to the use of a pharmaceutical composition as defined above in a method of providing symptomatic relief or treatment of pain, with or without fever, in particular that associated with inflammation such as that associated with trauma, osteoarthritis or rheumatoid arthritis, for example.

The invention also extends to the use of a combination of (i), (ii) and (iii) in the manufacture of a medicament for use in the symptomatic relief or treatment of pain, with or without fever.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

The pharmaceutical compositions of the invention are suitable for the symptomatic relief or treatment of pain with or without fever, in particular but not limited to that associated with inflammatory processes, such as trauma, osteoarthritis or rheumatoid arthritis, for example.

The first ingredient is an analgesic such as paracetamol (acetaminophen) or a pharmaceutically acceptable salt or derivative thereof. It has analgesic and antipyretic properties but limited or no anti-inflammatory action.

The daily dose of the paracetamol active ingredient is typically in the range of about 60 mg (children) to about 4000 mg (adults).

The second ingredient is a selective or specific COX-2 inhibitor such as meloxicam, celecoxib and rofecoxib, for example. These agents have anti-inflammatory and analgesic properties. Their ability to inhibit the action of COX-2 and not COX-1 has been shown to provide an enhanced safety profile for these compounds when compared to non-specific COX inhibitors.

In the case of meloxicam as active ingredient, the daily dose is typically in the range of about 3.75 mg to about 30 mg, preferably about 7.5 mg to about 15 mg.

The third ingredient is an opiate such as codeine, morphine, tramadol or fentanyl, for example. These compounds bind with specific receptors at many sites within the central nervous system to alter processes affecting both the perception of pain and the emotional response to pain.

The daily dose of the opiate, in the case of codeine phosphate, is 10 mg to 360 mg.

A pharmaceutical composition comprising a combination of an analgesic, an opiate and a selective or specific COX-2 inhibitor includes a pharmaceutically acceptable carrier and may include other necessary non-active excipients such as, for example, sorbitol, sucrose, saccharin, starch, lactose, guar gum, xanthan gum, magnesium stearate, bees wax, talc, methylcellulose, dextrin or povidone. The pharmaceutical composition may be provided in any appropriate dosage form such as, for example, tablets, capsules, granules, suspensions, solutions or other liquid forms, and is intended for oral, rectal or intravenous administration.

The dosage form will typically be administered to a patient from 2 to 4 times per day.

Although the active ingredients would typically be administered at currently accepted therapeutic doses, it is envisaged that one or more actives could be administered at lower than currently recognized doses while still providing effective pain relief/treatment.

DATED THIS 19th DAY OF FEBRUARY 2002

SPOOR & FISHER

APPLICANT'S PATENT ATTORNEYS